II. LISTING OF THE CLAIMS

The following listing of the claims replaces all prior versions, listing and amendments to the claims.

- 1. (Currently Amended) A method for reconstituting IKK in yeast comprising the steps of:
 - a. subcloning an IKK subunit genes gene into yeast expression vectors;
 - b. transforming said yeast expression vectors into yeast;
 - c. growing said yeast in a selective liquid media; and
 - d. controllably inducing the expression of said IKK subunits subunit by means of inducible promoters a promoter.
- 2. (Currently Amended) The method of claim 1, further comprising the steps of:
 - a. lysing said yeast;
 - b. extracting said an IKK protein produced by said IKK subunit gene; and
 - c. purifying said IKK protein.
- 3. (Currently Amended) The method of claim 1, wherein said yeast expression vectors contain a selection further comprise a gene encoding a selectable marker.
- 4. (Currently Amended) The method of claim 3, wherein said selection selectable marker gene encodes is leucine, histidine, tryptophan, or uracil.
- 5. (Currently Amended) The method of claim 1, wherein said yeast expression vectors contain-IKK subunits further comprise a polynucleotide encoding a tag.
- 6. (Currently Amended) The method of claim 1, wherein said tag is myc, HA, or FLAG or 6his.
- 7. (Currently Amended) The method of claim 1, wherein said yeast expression vectors contain an said promoter is an inducible promoter or a constitutive promoter.
- 8. (Previously Amended) The method of claim 7, wherein said inducible promoter is methionine or galactose.

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- 9. (Previously Amended) The method of claim 7, wherein said constitutive promoter is alcohol dehydrogenase.
- 10. (Original Claim) The method of claim 1, wherein said IKK subunit is IKKα.
- 11. (Original Claim) The method of claim 1, wherein said IKK subunit is IKK β .
- 12. (Original Claim) The method of claim 1, wherein said IKK subunit is IKKy.
- 13. (Currently Amended) The method of claim 1, wherein said IKK subunit comprises one or more of a combination of IKK α , IKK β , and IKK γ .
- 14. (Currently Amended) The method of claim 10, 11 or 13 wherein said <u>IKK subunit is</u>

 <u>IKKa and IKK\$ subunits are</u> subcloned into pESC ura or pESC trp vectors wherein a galactose promoter region is replaced with a met promoter from a leu(met) vector.
- 15. (Currently Amended) The method of claim 12 or 13, wherein said <u>IKKγIKK</u> subunit expression is subclosed into said leu(met) vector regulated by said promoter.
- 16. (Currently Amended) The method of claim 12 or 13, wherein said <u>IKKγIKK</u> subunit <u>expression</u> is subcloned into the <u>pES 86(+)</u> <u>a pESC</u> expression vector wherein constitutive <u>and said IKK</u> expression is induced under the alcehol dehydrogenase by a promoter.
- 17. (Original) The method of claim 1, wherein said yeast is Saccharomyces cerevisiae.
- 18. (Original) The method of claim 1, wherein said IKK is mammalian IKK.
- 19. (Previously Amended) The method of claim 18, wherein said mammalian IKK is human IKK.
- 20. (Original) The method of claim 1, wherein said vectors are plasmids, small yeast chromosomes or cosmids.
- 21. (Original) The method of claim 1, wherein said selective liquid media is an non-inducing drop-out media.
- 21. (Currently Amended) The method of claim 1, wherein said purified IKK protein-gene encodes wild-type IKK is substantially homologous to IKK isolated from wild-type cells.

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- 22. (Currently Amended) The method of claim 1, wherein said purified IKK gene protein is mutated.
- 23. (Original) A heterologously expressed IKK complex, wherein said IKK is expressed by yeast.
- 24. (Withdrawn) The composition of claim 24, wherein said IKK complex is comprised of IKK α , IKK β , and IKK γ subunits.
- 25. (Withdrawn) The composition of claim 24, wherein said IKK complex is produced by the method of claim 1.
- 26. (Withdrawn) A heterologously expressed IKK complex, wherein said IKKγ protein subunit regulates phosphorylation of serine residues in the activation of T loop kinase domain of IKK catalytic subunits.
- 27. (Withdrawn) The method of claim 27, wherein said IKK complex is activated by the dephosphorylation of γBD serines.
- 28. (Withdrawn) A yeast cell containing an expressible copy of a gene encoding a subunit of IKK.
- 29. (Withdrawn and Previously Amended) The yeast cell of claim 29 which is transformed with a yeast expression vector which contains the expressible copy of the gene encoding IKK α , IKK β , or IKK γ .
- 30. (Withdrawn and Previously Amended) The yeast cell of claim 29 which is transformed by the method of claim 1.
- 31. (Withdrawn) A method for identifying upstream regulators of IKK complex, comprising the steps of:
 - a. mutating the genes of one or more said IKK subunits;
 - b. subcloning genes for IKK subunits into yeast expression vectors;
 - c. transforming said yeast expression vectors into yeast;

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- d. growing said yeast in a selective liquid media;
- e. controllably inducing the expression of said IKK subunits by means of inducible promoters;
- f. lysing said yeast;
- g. extracting said IKK protein;
- h. purifying said IKK protein; and
- comparing kinase activity of said IKK protein with wild type IKK.
- 32. (Withdrawn) The method of claim 32, wherein said mutation is on a binding domain.
- 33. (Withdrawn and Previously Amended) The method of claim 33, wherein said mutation mimics the biochemical characteristics of said binding site when bound.
- 34. (Withdrawn and Previously Amended) The method of claim 33, wherein said mutation prevents binding at said domain site.
- 35. (Withdrawn) The method of claim 32, wherein said mutation changes serines to alanines.
- 36. (Withdrawn) The method of claim 32, wherein said mutation changes serines to glutamic acid.
- 37. (Withdrawn) A method for assaying IKK activity in situ in yeast comprising the steps of:
 - a. subcloning genes for IKK subunits into first yeast expression vectors;
 - b. transforming said first yeast expression vectors into yeast;
 - subcloning HeLa cell cDNA into second yeast expression vectors;
 - d. transforming said yeast expression vectors into said yeast;
 - e. replica plating said yeast;
 - f. growing said yeast on membranes on selective non-inducing medium
 - g. inducing said yeast to produce IKK protein;
 - h. fixing said IKK protein;

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- 39. (Withdrawn and Previously Amended) The method of claim 38, further comprising the step of sequencing said positive clones.
- 40. (Withdrawn and Previously Amended) The method of claim 38, further comprising the steps of:
 - a. transforming said positive clone into yeast;
 - b. growing said yeast in a selective liquid media;
 - c. controllably inducing the expression of said clones by means of inducible promoters.
- 41. (Withdrawn and Previously Amended) The method of claim 0, further comprising the steps of:
 - a. transforming said positive clone into yeast;
 - growing said yeast in a selective liquid media;
 - c. controllably inducing the expression of said clones by means of inducible promoters.

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